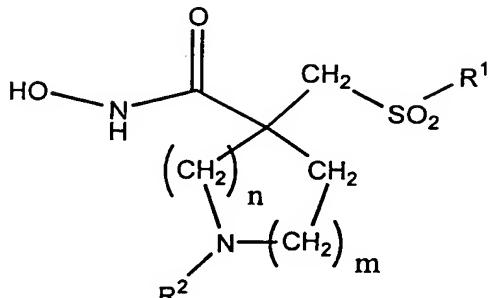


WHAT IS CLAIMED IS:

1. A compound corresponding to Formula I:



I

5

wherein

m is zero, 1, 2 or 3;

n is zero, 1, or 2, and the sum of m plus n
is 1, 2 or 3;

10 R² is hydrido, C₁-C₈ hydrocarbyl, C₁-C₆
hydrocarbyloxycarbonyl C₁-C₄ hydrocarbyl, aryl C₁-C₄
hydrocarbyl, heteroaryl C₁-C₄ hydrocarbyl, aryloxy
C₁-C₄ hydrocarbyl, or heteroarylcxy C₁-C₄
hydrocarbyl; and

15 R¹ is a substituent containing a 5- or 6-
membered cyclohydrocarbyl, heterocyclo, aryl or
heteroaryl radical bonded directly to the depicted
SO₂-group and having a length greater than about that
of a hexyl group and less than about that of an

20 eicosyl group, said R¹ defining a three-dimensional
volume, when rotated about an axis drawn through the
SO₂-bonded 1-position and the 4-position of a 6-
membered ring radical or drawn through the SO₂-bonded
1-position and the center of 3,4-bond of a 5-membered

ring radical, whose widest dimension in a direction transverse to the axis of rotation is about that of one furanyl ring to about that of two phenyl rings.

5 2. The compound according to claim 1 wherein said 5- or 6-membered cyclohydrocarbyl, heterocyclo, aryl or heteroaryl radical of R¹ is substituted with a substituent, R³, that has a chain length of 3 to about 14 carbon atoms.

10

3. The compound according to claim 2 wherein said R³ substituent is selected from the group consisting of a phenyl group, a phenoxy group, a thiophenoxy group, an anilino group, a phenylazo group, a phenylureido, a benzamido], a nicotinamido, an isonicotinamido, a picolinamido group, a heterocyclo, heterocyclohydrocarbyl, arylheterocyclohydrocarbyl, arylhydrocarbyl, heteroarylhydrocarbyl,
15 heteroarylheterocyclohydrocarbyl, arylhydrocarbyloxyhydrocarbyl, aryloxyhydrocarbyl, hydrocarboylhydrocarbyl, arylhydrocarboylhydrocarbyl, arylcarbonylhydrocarbyl, arylazoaryl, arylhydrazinoaryl, hydrocarbylthiohydrocarbyl,
20 hydrocarbylthioaryl, arylthiohydrocarbyl, heteroarylthiohydrocarbyl, hydrocarbylthioarylhydrocarbyl, arylhydrocarbylthiohydrocarbyl, arylhydrocarbylthioaryl, arylhydrocarbylamino,
25 heteroarylhydrocarbylamino, and a heteroarylthio group.
30

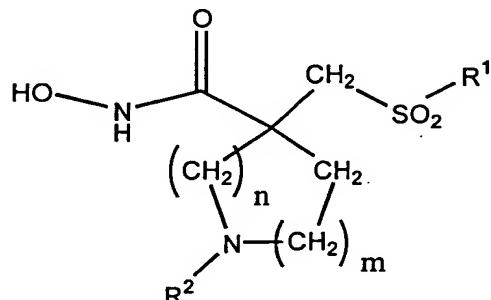
4. The compound according to claim 3

wherein said R³ substituent is itself substituted by
one or more substituents selected from the group
consisting of a halogen, hydrocarbyl, hydrocarbyloxy,
5 nitro, cyano, perfluorohydrocarbyl,
trifluoromethylhydrocarbyl, hydroxy, mercapto,
hydroxycarbonyl, aryloxy, arylthio, arylamino,
arylhydrocarbyl, aryl, heteroaryloxy, heteroarylthio,
heteroarylamino, heteroarylhydrocarbyl,
10 hydrocarbyloxycarbonylhydrocarbyl, heterocyclooxy,
hydroxycarbonylhydrocarbyl, heterocyclothio,
heterocycloamino, cyclohydrocarbyloxy,
cyclohydrocarbylthio, cyclohydrocarbylamino,
heteroarylhydrocarbyloxy, heteroarylhydrocarbylthio,
15 heteroarylhydrocarbylamino, arylhydrocarbyloxy,
arylhydrocarbylthio, arylhydrocarbylamino,
heterocyclic, heteroaryl, hydroxycarbonyl-
hydrocarbyloxy, alkoxy carbonylalkoxy, hydrocarbyloyl,
arylcarbonyl, arylhydrocarbyloyl, hydrocarboyoxy,
20 arylhydrocarboyoxy, hydroxyhydrocarbyl,
hydroxyhydrocarbyloxy, hydrocarbylthio,
hydrocarbyloxyhydrocarbylthio,
hydrocarbyloxy carbonyl,
hydroxycarbonylhydrocarbyloxy, hydrocarbyloxy-
25 carbonylhydrocarbyl, hydrocarbylhydroxycarbonyl-
hydrocarbylthio,
hydrocarbyloxy carbonylhydrocarbyloxy,
hydrocarbyloxy carbonylhydrocarbylthio, amino,
hydrocarbyl carbonylamino, aryl carbonylamino,
30 cyclohydrocarbyl carbonylamino,
heterocyclohydrocarbyl carbonylamino,
arylhydrocarbyl carbonylamino,
heteroaryl carbonylamino,

heteroarylhydrocarbylcarbonylamino,
heterocyclohydrocarbyloxy, hydrocarbysulfonylamino,
arylsulfonylamino, arylhydrocarbysulfonylamino,
heteroarylsulfonylamino, heteroarylhydrocarbyl-
5 sulfonylamino, cyclohydrocarbysulfonylamino,
heterocyclohydrocarbysulfonylamino and N-
monosubstituted or N,N-disubstituted aminohydrocarbyl
group, wherein the substituent(s) on the nitrogen are
selected from the group consisting of hydrocarbyl,
10 aryl, arylhydrocarbyl, cyclohydrocarbyl,
arylhydrocarbyloxycarbonyl, hydrocarbyloxycarbonyl,
and hydrocarboyl, or wherein the nitrogen and two
substituents attached thereto form a 5- to 8-membered
heterocyclic or heteroaryl ring group.

15

5. A compound corresponding to Formula I:



20

wherein

m is zero, 1, 2 or 3;

n is zero, 1, or 2, and the sum of m plus n
is 1, 2 or 3;

R² is hydrido, C₁-C₈ hydrocarbyl, C₁-C₆ hydrocarbyloxycarbonyl C₁-C₄ hydrocarbyl, aryl C₁-C₄ hydrocarbyl, heteroaryl C₁-C₄ hydrocarbyl, aryloxy C₁-C₄ hydrocarbyl, or heteroaryloxy C₁-C₄ hydrocarbyl; and

R¹ is a substituent containing a 5- or 6-membered cyclohydrocarbyl, heterocyclo, aryl or heteroaryl radical bonded directly to the depicted SO₂-group that is itself substituted at its own 4-position when a 6-membered ring and at its own 3- or 4-position when a 5-membered ring with a substituent R³ selected from the group consisting of one other single-ringed cyclohydrocarbyl, heterocyclo, aryl or heteroaryl group, a C₃-C₁₄ hydrocarbyl group, a C₂-C₁₄ hydrocarbyloxy group, a phenoxy group, a thiophenoxy group, a 4-thiopyridyl group, a phenylazo group, a phenylureido group, a nicotinamido group, an isonicotinamido group, a picolinamido group, an anilino group and a benzamido group.

20

6. The compound according to claim 5 wherein said R¹ substituent is PhR³ in which Ph is phenyl substituted with R³ at the 4-position, and R³ is a phenyl, phenoxy, thiophenoxy, phenylazo, benzamido, anilino, nicotinamido, isonicotinamido, picolinamido or phenylureido group

7. The compound according to claim 5 wherein said R¹ substituent is PhR³ in which Ph is phenyl substituted with R³ at the 4-position, and said R³ is a phenyl, phenoxy, anilino or thiophenoxy

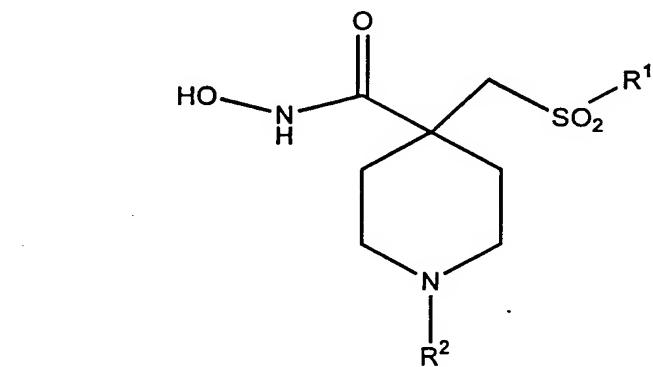
group that is optionally substituted at the meta- or para-position or both with a moiety that is selected from the group consisting of a halogen, a C₁-C₉ hydrocarbyloxy group, a C₁-C₁₀ hydrocarbyl group, a 5 di- C₁-C₉ hydrocarbylamino group, a carboxyl C₁-C₈ hydrocarbyl group, a C₁-C₄ hydrocarbyloxy carbonyl C₁-C₄ hydrocarbyl group, a C₁-C₄ hydrocarbyloxycarbonyl C₁-C₄ hydrocarbyl group and a carboxamido C₁-C₈ hydrocarbyl group, or is 10 substituted at the meta- and para-positions by two methyl groups or by a methylenedioxy group.

8. The compound according to claim 5 wherein m is 2 and n is zero.

15

9. The compound according to claim 5 wherein m is 1 and n is 2.

20 II:



II

wherein

R² is hydrido, C₁-C₈ hydrocarbyl, C₁-C₆ hydrocarbyloxycarbonyl C₁-C₄ hydrocarbyl, aryl C₁-C₄ hydrocarbyl, heteroaryl C₁-C₄ hydrocarbyl, aryloxy C₁-C₄ hydrocarbyl, or heteroaryloxy C₁-C₄ hydrocarbyl; and

R¹ is a substituent containing a 5- or 6-membered cyclohydrocarbyl, heterocyclo, aryl or heteroaryl radical bonded directly to the depicted SO₂-group that is itself substituted at its own 4-position when a 6-membered ring and at its own 3- or 4-position when a 5-membered ring with a substituent R³ selected from the group consisting of one other single-ringed cyclohydrocarbyl, heterocyclo, aryl or heteroaryl group, a C₃-C₁₄ hydrocarbyl group, a C₂-C₁₄ hydrocarbyloxy group, a phenoxy group, a thiophenoxy group, a 4-thiopyridyl group, a phenylazo group, a phenylureido group, a nicotinamido group, an isonicotinamido group, a picolinamido group, an anilino group and a benzamido group.

11. The compound according to claim 10 wherein R¹ is a single-ringed aryl or heteroaryl group that is 5- or 6-membered, and is itself substituted at its own 4-position when a 6-membered ring and at its own 3- or 4-position when a 5-membered ring with a substituent selected from the group consisting of one other single-ringed aryl or heteroaryl group, a C₆-C₁₄ hydrocarbyl group, a C₆-C₁₄ hydrocarbyloxy group, a phenoxy group, a thiophenoxy group, a 4-thiopyridyl group, a phenylazo group a phenylureido group and a benzamido group.

12. The compound according to claim 10
wherein said R¹ radical is PhR³ in which Ph is phenyl
substituted with R³ at the 4-position, and R³ is a
5 phenyl, phenoxy, thiophenoxy, phenylazo, benzamido,
nicotinamido, isonicotinamido, picolinamido or
phenylureido group

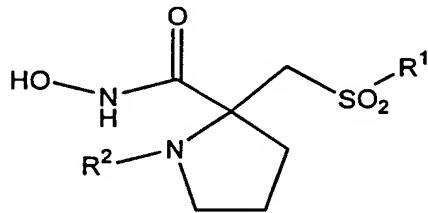
13. The compound according to claim 10
10 wherein said R¹ radical is PhR³ in which Ph is phenyl
substituted with R³ at the 4-position, and said R³ is
a phenyl, phenoxy or thiophenoxy group that is
optionally substituted at the meta- or para-position
or both with a moiety that is selected from the group
15 consisting of a halogen, a C₁-C₉ hydrocarbyloxy
group, a C₁-C₁₀ hydrocarbyl group, a di- C₁-C₉
hydrocarbylamino group, a carboxyl C₁-C₈ hydrocarbyl
group, a C₁-C₄ hydrocarbyloxy carbonyl C₁-C₄
hydrocarbyl group, a C₁-C₄ hydrocarbyloxycarbonyl C₁-
20 C₄ hydrocarbyl group and a carboxamido C₁-C₈
hydrocarbyl group, or is substituted at the meta- and
para-positions by two methyl groups or by a
methylenedioxy group.

25 14. The compound according to claim 10
wherein said R¹ radical is PhR³ in which Ph is phenyl
substituted with R³ at the 4-position, and said R³
substituent is an benzamido, nicotinamido, an
anilino, isonicotinamido, picolinamido or
30 phenylureido group in which said R³ substituent is
optionally substituted at its own meta- or para-

position or both with a moiety selected from the group consisting of a halogen, a nitro, a C₁-C₈ hydrocarbyl, a C₁-C₇ hydrocarbyloxy, an amino and an amino-C₂-C₄-hydroxyalkyl group.

5

15. A compound corresponding to Formula IV:



10

IV

wherein:

R² is hydrido, C₁-C₈ hydrocarbyl, C₁-C₆ hydrocarbyloxycarbonyl C₁-C₄ hydrocarbyl, aryl C₁-C₄ hydrocarbyl, heteroaryl C₁-C₄ hydrocarbyl, aryloxy C₁-C₄ hydrocarbyl, or heteroaryloxy C₁-C₄ hydrocarbyl; and

R¹ is a substituent containing a single aryl or heteroaryl radical bonded directly to the 20 depicted SO₂-group that is itself substituted at its own 4-position when a 6-membered ring and at its own 3- or 4-position when a 5-membered ring with a substituent selected from the group consisting of one other single-ringed aryl or heteroaryl group, a C₃-C₁₄ hydrocarbyl group, a C₂-C₁₄ hydrocarbyloxy group, a phenoxy group, a thiophenoxy group, a 4-thiopyridyl

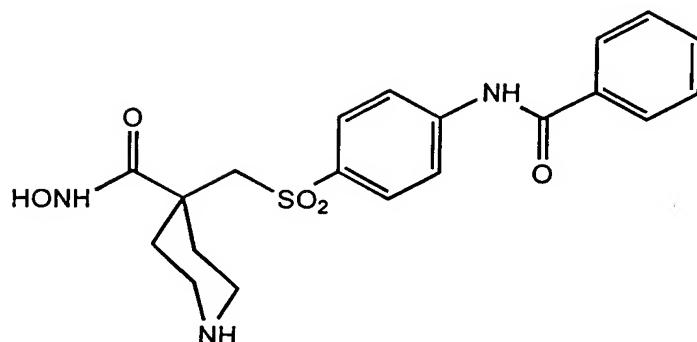
group, a phenylazo group, a phenylureido group, a nicotinamido group, an isonicotinamido group, a picolinamido group, an anilino group and a benzamido group.

5

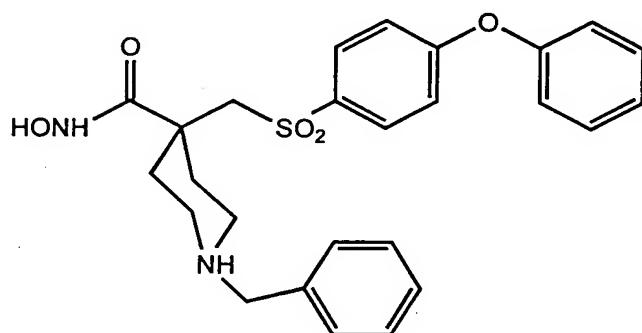
16. The compound according to claim 15 wherein said R¹ radical is PhR³ in which Ph is phenyl substituted with R³ at the 4-position, and R³ is a phenyl, phenoxy, thiophenoxy, anilino, phenylazo, 10 benzamido, nicotinamido, isonicotinamido, picolinamido or phenylureido group.

17. The compound according to claim 15 wherein said R¹ radical is PhR³ in which Ph is phenyl 15 substituted with R³ at the 4-position, and said R³ is a phenyl, phenoxy, anilino, thiophenoxy, benzamido, nicotinamido, isonicotinamido, picolinamido or phenylureido group that is optionally substituted at the meta- or para-position or both with a moiety that 20 is selected from the group consisting of a halogen, a C₁-C₉ hydrocarbyloxy group, a C₁-C₁₀ hydrocarbyl group, a di- C₁-C₉ hydrocarbylamino group, a carboxyl C₁-C₈ hydrocarbyl group, a C₁-C₄ hydrocarbyloxy carbonyl C₁-C₄ hydrocarbyl group, a C₁-C₄ 25 hydrocarbyloxycarbonyl C₁-C₄ hydrocarbyl group and a carboxamido C₁-C₈ hydrocarbyl group, or is substituted at the meta- and para-positions by two methyl groups or by a methylenedioxy group.

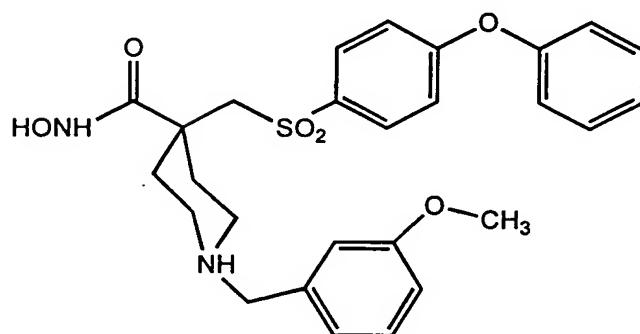
18. A compound corresponding in structure to the formula



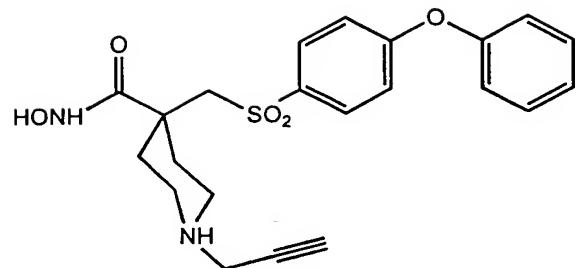
5 19. A compound corresponding in structure to the formula



10 20. A compound corresponding in structure to the formula

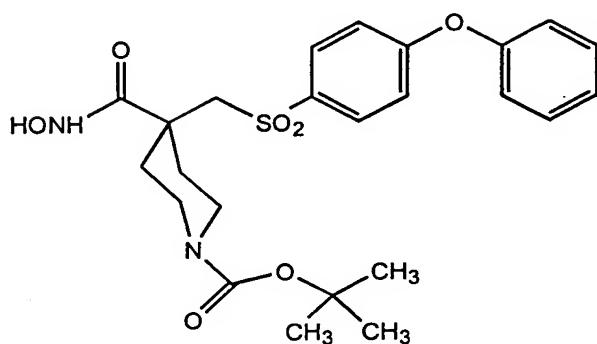


21. A compound corresponding in structure to the formula



22. A compound corresponding in structure
to the formula

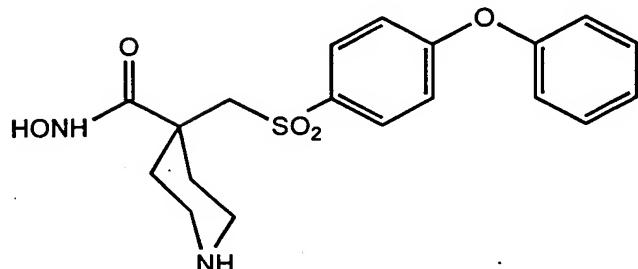
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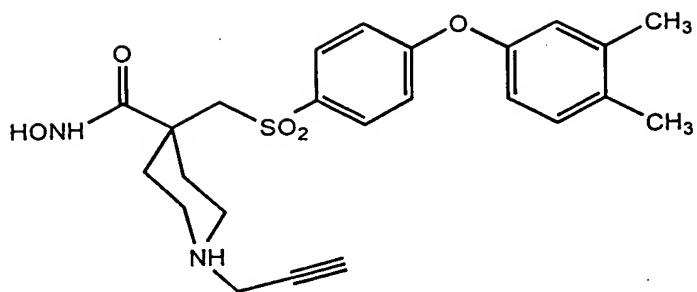
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23. A compound corresponding in structure
to the formula

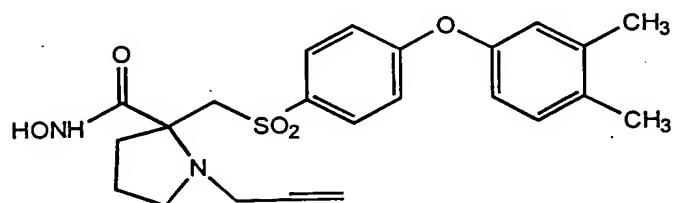
15



24. A compound corresponding in structure
20 to the formula

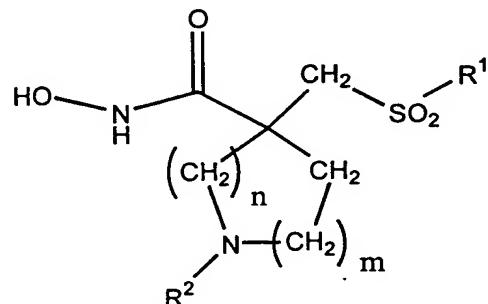


25. A compound corresponding in structure to the formula



5

26. A process for treating a host mammal having a condition associated with pathological matrix metalloprotease activity that comprises
10 administering a compound corresponding in structure to Formula I in an MMP enzyme-inhibiting effective amount to a mammalian host having such a condition:



I

15

wherein

m is zero, 1, 2 or 3;

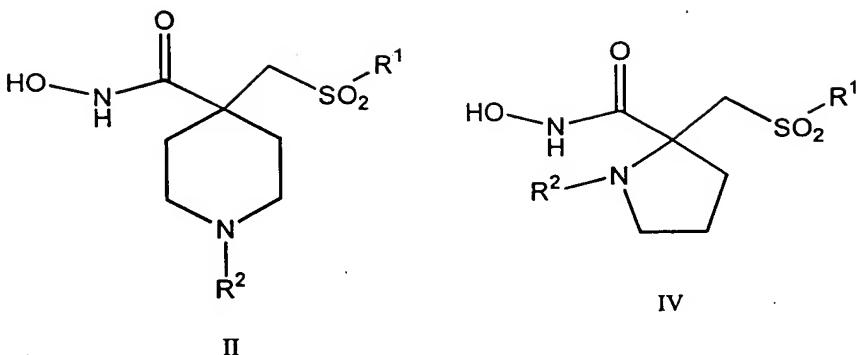
n is zero, 1, or 2, and the sum of m plus n
is 1, 2 or 3;

R² is hydrido, C₁-C₈ hydrocarbyl, C₁-C₆

5 hydrocarbyloxycarbonyl C₁-C₄ hydrocarbyl, aryl C₁-C₄
hydrocarbyl, heteroaryl C₁-C₄ hydrocarbyl, aryloxy
C₁-C₄ hydrocarbyl, or heteroaryloxy C₁-C₄
hydrocarbyl; and

10 R¹ is a substituent containing a 5- or 6-
membered cyclohydrocarbyl, heterocyclo, aryl or
heteroaryl radical bonded directly to the depicted
SO₂-group and having a length greater than about that
of a hexyl group and less than about that of an
eicosyl group, said R¹ defining a three-dimensional
15 volume, when rotated about an axis drawn through the
SO₂-bonded 1-position and the 4-position of a 6-
membered ring radical or drawn through the SO₂-bonded
1-position and the center of 3,4-bond of a 5-membered
ring radical, whose widest dimension in a direction
20 transverse to the axis of rotation is about that of
one furanyl ring to about that of two phenyl rings.

27. The process according to claim 26
wherein said compound corresponds in structure to
25 either Formula II or Formula IV:



28. The process according to claim 27
wherein R¹ is a single-ringed aryl or heteroaryl
5 group that is 5- or 6-membered, and is itself
substituted at its own 4-position when a 6-membered
ring and at its own 3- or 4-position when a
5-membered ring with a substituent selected from the
group consisting of one other single-ringed aryl or
10 heteroaryl group, a C₃-C₁₄ hydrocarbyl group, a C₂-
C₁₄ hydrocarbyloxy group, a phenoxy group, a
thiophenoxy group, an anilino group, a 4-thiopyridyl
group, a phenylazo group a phenylureido group, a
nicotinamido group, an isonicotinamido group, a
15 picolinamido group and a benzamido group.

29. The process according to claim 27
wherein said R¹ radical is PhR³ in which Ph is phenyl
substituted with R³ at the 4-position, and R³ is a
20 phenyl, phenoxy, anilino, thiophenoxy, phenylazo,
benzamido, nicotinamido, isonicotinamido,
picolinamido or phenylureido group.

30. The process according to claim 29
25 wherein said R¹ radical is PhR³ in which Ph is phenyl
substituted with R³ at the 4-position, and said R³ is

a phenyl, phenoxy, anilino or thiophenoxy group that is optionally substituted at the meta- or para-position or both with a moiety that is selected from the group consisting of a halogen, a C₁-C₉ hydrocarbyloxy group, a C₁-C₁₀ hydrocarbyl group, a di-C₁-C₉ hydrocarbylamino group, a carboxyl C₁-C₈ hydrocarbyl group, a C₁-C₄ hydrocarbyloxy carbonyl C₁-C₄ hydrocarbyl group, a C₁-C₄ hydrocarbyloxycarbonyl C₁-C₄ hydrocarbyl group and a carboxamido C₁-C₈ hydrocarbyl group, or is substituted at the meta- and para-positions by two methyl groups or by a methylenedioxy group.

31. The process according to claim 26
15 wherein said compound is administered a plurality of times.